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Attorney Docket No. 1567.1059

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Patent Application of:

Jea-Woan LEE

Application No.: 10/691,476

Group Art Unit:

Confirmation No.: 5031

Filed: October 23, 2003

Examiner:

For: **NEGATIVE ELECTRODE FOR LITHIUM BATTERY AND LITHIUM BATTERY  
COMPRISING SAME**

**SUBMISSION OF CERTIFIED COPY OF PRIOR FOREIGN  
APPLICATION IN ACCORDANCE  
WITH THE REQUIREMENTS OF 37 C.F.R. § 1.55**

Commissioner for Patents  
PO Box 1450  
Alexandria, VA 22313-1450

Sir:

In accordance with the provisions of 37 C.F.R. § 1.55, the applicant(s) submit(s) herewith  
a certified copy of the following foreign application:

Korean Patent Application No(s). 2002-65484

Filed: October 25, 2002

It is respectfully requested that the applicant(s) be given the benefit of the foreign filing  
date(s) as evidenced by the certified papers attached hereto, in accordance with the  
requirements of 35 U.S.C. § 119.

Respectfully submitted,

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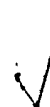
Submission of Certified Copy of Prior Foreign Application, Korean Priority Document 2002-65484, no fees required

APPLICANT(S): Jea-Woan LEE  
SERIAL NO:  
CONFIRMATION NO. 5031  
TITLE: NEGATIVE ELECTRODE FOR LITHIUM BATTERY AND LITHIUM BATTERY COMPRISING SAME  
FILING DATE: October 23, 2003  
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APPLICANT(S): Jea-Woan LEE  
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Attorney Docket No. 1567.1059

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of:

Jea-Woan LEE

Application No.: 10/691,476

Group Art Unit: 1745

Filed: October 23, 2003

Examiner: Laura S. Weiner

For: NEGATIVE ELECTRODE FOR LITHIUM BATTERY AND LITHIUM BATTERY  
COMPRISING SAME

**SUBMISSION OF VERIFIED TRANSLATION OF CERTIFIED COPY OF PRIOR FOREIGN  
APPLICATION**

Commissioner for Patents  
PO Box 1450  
Alexandria, VA 22313-1450

Sir:

Applicants submit herewith a translation of Korean Patent Application No. 2002-65484 claiming priority to October 25, 2002 and a statement from the translator.

If there are any fees associated with filing of this Submission, please charge the same to our Deposit Account No. 503333.

Respectfully submitted,

STEIN, MCEWEN & BUI, LLP

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## Verification Statement For Translation

I, YOM, Chol-jong, hereby declare that I am conversant in the Korean and the English languages and that I am the translator of the document attached and certify that to the best of my knowledge and belief the following is a true and correct English translation of the Korean Patent Application No. 2002-0065484 filed on October 25, 2002.

Signature : 염철종 YOM, Chol-jong  
Date : January 16, 2007



## 【ABSTRACT OF THE DISCLOSURE】

### 【ABSTRACT】

The present invention relates to a negative electrode for a lithium battery, and a lithium battery comprising the same.

Disclosed is a negative electrode for a lithium battery comprising a metallic lithium plate and a negative electrode tab attached to the surface of the metallic lithium plate, wherein an average surface roughness of the metallic lithium plate on the area attached to the negative tab is 0.1 to 5  $\mu\text{m}$ ; a negative electrode for a lithium battery comprising a metallic lithium plate and a negative electrode tab having a porosity of 50 to 100% and being attached to the metallic lithium plate; a negative electrode for a lithium battery comprising a metallic lithium plate and a negative electrode tab attached to both the upper and lower end surfaces of the metallic lithium plate; or a negative electrode for a lithium battery comprising a metallic lithium plate and a negative electrode tab attached to the surfaces of the metallic lithium plate, wherein the surface area of the negative electrode tab to be attached to the metallic lithium plate is 10% larger than the geographical area.

### 【REPRESENTATIVE DRAWINGS】

Fig. 1

### 【KEYWORD】

lithium battery, negative tab, surface roughness, porosity, foam

## **【SPECIFICATION】**

### **【TITLE】**

NEGATIVE ELECTRODE FOR LITHIUM BATTERY AND LITHIUM  
BATTERY COMPRISING SAME

### **【BRIEF DESCRIPTION OF THE DRAWINGS】**

A more complete appreciation of the invention, and many of the attendant advantages thereof, will be readily apparent as the same becomes better understood by reference to the following detailed description when considered in conjunction with the accompanying drawing, wherein:

FIG. 1 shows a cross-sectional view of a lithium battery.

- |                            |                            |
|----------------------------|----------------------------|
| 1: battery                 | 2: positive electrode      |
| 4: negative electrode      | 6: separator               |
| 8: electrode group         | 10: case                   |
| 12: battery cover          | 14: gasket                 |
| 16: positive electrode tab | 18: negative electrode tab |
| 20, 22: insulator          | 24: electrolyte            |

### **【DETAILED DESCRIPTION OF THE INVENTION】**

### **【OBJECT OF THE INVENTION】**

**【TECHNICAL FIELD OF THE INVENTION AND DESCRIPTION OF THE  
RELATED ART】**

The present invention relates to a negative electrode for a lithium battery and a lithium battery comprising the same, and more particularly, to a negative electrode for a lithium battery that is capable of enhancing capacity characteristics and decreasing the occurrence of a short therein, and a lithium battery comprising the same.

As portable electronic products have tended to become more and more miniature in size and lighter in weight, demands for development of batteries exhibiting higher performance and higher capacity have rapidly increased. Generally, batteries are divided into primary (non-rechargeable) and secondary (rechargeable) batteries, depending on their capability of being electrically recharged. The primary battery may include, for example, a manganese battery, an alkaline battery, a mercury battery, a silver oxide battery, and so on. The secondary battery may include, for example, a lead battery, a Ni-MH (nickel-metal hydride) battery, a sealed nickel-cadmium battery, a metallic lithium battery, a lithium ion battery, a lithium polymer battery, a lithium-sulfur battery, and so on.

Batteries are electrochemical devices that convert chemical energy into electrical energy by electrochemical oxidation and reduction reactions between a positive and a negative electrode. The active materials participating in the electrochemical reaction between these two electrodes influence the confidence and the performance factors of the battery such as capacity, cycle life, and voltage.

Lithium is an attractive material among currently used active materials since lithium has a high electric capacity per unit of weight and high electronegativity, which is anticipated to impart a high capacity and a high voltage to a battery. In a case of employing metallic lithium as a negative active material, the metallic lithium can be used both as the active material and as a current collector at the same time. The metallic lithium plate is thus used as a negative electrode plate by itself, without adding a current collector.

FIG. 1 shows a structure of a non-aqueous lithium battery 1. The battery is fabricated by the following steps of interposing a separator 6 between a positive electrode 2 and a negative electrode 4, winding them to form an electrode group 8, and inserting the electrode group into a case 10. The upper side of the battery case 10 is then sealed with a battery cover 12 and a gasket 14, and a safety vent may be installed in the battery cover 12 to permit the escape of gases. The outer surface of the battery cover 12 acts as a positive electrode pole, while the outer surface of the case 10 acts as a negative electrode pole. The positive electrode tab 16 and the negative electrode tab 18 are connected so that the electrodes are associated with the poles. Insulators 20, 22 are placed inside the battery to prevent the occurrence of a short, and electrolyte 24 is injected therein prior to sealing the battery by clamping the cover 12 on the case 10.

When the negative electrode plate is a metallic lithium negative electrode



and the battery case is made of a metallic material, the metallic lithium negative electrode would be directly connected to the battery case so that the electricity is conducted therebetween. This, however, could cause a problem in that the electric conductivity is degenerated between the outer surface of the battery case and the metallic lithium since the metallic lithium tends to react with electrolytes. Meanwhile, when the battery case is not made of metallic materials, it is necessary to take the pole outside of the battery, and accordingly, the negative electrode tab should be made of materials that are not dissolved and eluted in the electrolyte.

Nowadays, electric devices such as portable phones require a pouch-type battery since they are lighter in weight, higher in capacity, and rectangular in shape. The lithium is known to have a high capacity per unit of weight, so that it is becoming attractive as a negative active material. In addition to this tendency, methods for electrically connecting with the battery pole in a case of employing the metallic lithium as a negative electrode are being vigorously studied.

Japanese Patent Laid-Open Publication No. P5-251073 discloses a method of preventing the edge of a nickel tab from damaging the separator, and a method of decreasing occurrence of a short by covering the nickel tab with the metallic lithium in a manner such that the nickel tab is stacked on the lithium foil and the lithium is further stacked thereon. The method, however, causes a problem in that

the effective capacity of the battery is decreased by as much as the space occupied by the lithium covering the nickel tab.

#### **【TECHNICAL OBJECT TO BE ACCOMPLISHED OF THE INVENTION】**

The object of present invention is to provide a negative electrode for a lithium battery having a high capacity as well as an excellent attaching strength between a metallic lithium negative electrode and a negative electrode tab.

Another object of present invention is to provide a method of fabricating a negative electrode for a lithium battery having a high capacity and an excellent attaching strength between a metallic lithium negative electrode and a negative electrode tab.

#### **【DETAILED DESCRIPTION OF THE INVENTION】**

In order achieve the objects, the present invention provides a negative electrode for a lithium battery, comprising a metallic lithium plate and a negative electrode tab attached to the surface of the metallic lithium plate, wherein an average surface roughness (Ra) of the metallic lithium plate on the area attached to the negative electrode tab is 0.1 to 5  $\mu\text{m}$ .

The present invention also provides a negative electrode for a lithium battery, comprising a metallic lithium plate and a negative electrode tab attached to the metallic lithium plate, wherein the negative electrode tab has a porosity of 50 to

100%.

The present invention further provides a negative electrode for a lithium battery, comprising a metallic lithium plate and a negative electrode tab attached to both the upper and lower end surfaces of the metallic lithium plate.

The present invention further provides a negative electrode for a lithium battery, comprising a metallic lithium plate and a negative electrode tab attached to the surface of metallic lithium plate, wherein the surface of negative electrode tab that is attached to the metallic lithium plate has a surface area of 10% larger than a geographical area.

The present invention further provides a method of fabricating a negative electrode for a lithium battery, comprising the steps of brushing the surface area of a metallic lithium plate to be attached to a negative electrode tab so that the average surface roughness (Ra) of the surface area is 0.1 to 5  $\mu\text{m}$ , and pressing the negative electrode tab onto the metallic lithium plate and attaching the negative electrode tab with the metallic lithium plate.

According to the first embodiment of the present invention, a negative electrode for a lithium secondary battery comprises a metallic lithium plate and a negative electrode tab attached to the surface of the metallic lithium plate, wherein an average surface roughness (Ra) of the metallic lithium plate at the area

attached to the negative tab is 0.1 to 5  $\mu\text{m}$ .

The metallic lithium plate is preferably a metallic lithium foil or metallic lithium coated on a conductive substrate. The representative example of the conductive substrate includes a metal foil, a metal film, a conductive polymer film, and a polymer film deposited with a metal. The metal foil and the metal film may be composed of copper or nickel. The term "polymer film deposited with a metal" means that any metal such as copper or nickel is deposited on a polymer film. The polymer film may be composed of polyacetylene, polypyrrole, polyaniline, polythiophene, poly(p-phenylene), poly(phenylene vinylene), polyazulene, poly(perinaphthalene), poly(naphthalene-2,6-diyl), polyacene, and so on.

The negative electrode tab is preferably made of a metal plate or a metal foam having a thickness of 10 to 50  $\mu\text{m}$ . The metal plate and the metal foam may be composed of nickel, copper, iron, stainless steel, and so on. The average surface roughness (Ra) is preferably in a range of 0.1 to 5  $\mu\text{m}$ , and more preferably 0.3 to 0.6  $\mu\text{m}$ . In a case of when the roughness is less than 0.1  $\mu\text{m}$ , the negative electrode tab cannot be firmly attached to the plate, while in a case of when the roughness is more than 5  $\mu\text{m}$ , it may cause a problem in that the electrode plate

gets damaged or the tab gets broken and disconnected during the brushing process.

A sheet-type negative electrode tab is stacked on the surface of the metallic lithium plate which has the aforementioned range of roughness, and then the tab is pressed to impart a firm attachment to the plate. However, the method of attaching the negative electrode tab to the negative electrode plate is not limited to pressing.

According to the second embodiment of the present invention, the negative electrode for a lithium battery comprises a metallic lithium plate and a negative electrode tab attached to the metallic lithium plate, wherein the negative electrode tab has a porosity of 50 to 100%.

The metallic lithium plate is a metallic lithium foil or metallic lithium coated on a conductive substrate, which are identical to those in the first embodiment.

The negative electrode tab is in a form of foam, and has a porosity of 50 to 100% and preferably 80 to 95%. When the negative electrode tab is made of a foam having a porosity within the range, the welding of the negative electrode tab to the metallic lithium plate is effectively performed since the lithium is pressed and incorporated into the void of the foam or melted and coagulated into the void of the foam.

The foam negative electrode tab is stacked on the surface of the metallic

lithium plate, followed by pressing them to perform the firm attachment of the negative electrode tab to the negative electrode plate.

According to the third embodiment, the negative electrode for a lithium battery comprises a metallic lithium plate, and a negative electrode tab attached to both upper and lower end surfaces of the metallic lithium plate.

The metallic lithium plate is preferably a metallic lithium foil or metallic lithium coated on a conductive substrate, which are identical to those in the first embodiment.

The negative electrode tab may be a metal foil or a metal foam. The negative electrode tab may be composed of, but is not limited to, nickel, copper, iron, stainless steel, and so on. An upper tab and a lower tab are placed parallel with each other on both end surfaces of the metallic lithium plate, and the upper tab is welded to the lithium and the lower tab is welded to the lithium.

According to the fourth embodiment of the present invention, a negative electrode for a lithium battery comprises a metallic lithium plate and a negative electrode tab attached to the surface of metallic lithium plate, wherein the surface of negative electrode tab that is attached to the metallic lithium plate has a surface area of 10% larger than a geographical area.

The metallic lithium plate is preferably a metallic lithium foil or metallic lithium coated on a conductive substrate, which are identical to those in the first

embodiment.

The negative electrode tab may be a metal foil or a metal foam. The negative electrode tab may be composed of, but is not limited to, nickel, copper, iron, stainless steel, and so on. The surface area of the negative electrode tab contacted with the metallic lithium plate is increased by 10%, preferably by 50 to 100% compared to the geographical area. The term "geographical area" is intended to mean the surface area supposing the surface is complete flat. In order to increase the contact of the negative electrode tab, the average surface roughness must be controlled. The average surface roughness of the area of negative electrode tab contacted with the metallic lithium plate is preferably 0.1 to 5  $\mu\text{m}$ , and more preferably 0.3 to 0.6  $\mu\text{m}$ . In a case of when the roughness is less than 0.1  $\mu\text{m}$ , the tab cannot be firmly attached to the metallic lithium plate, and in a case of when the roughness is more than 5  $\mu\text{m}$ , the tab can be more easily broken or disconnected.

When the negative electrode tab is firmly attached to the metallic lithium plate, it is possible to provide a battery having a high capacity since the internal resistance is decreased upon the charge and discharge of the battery, to suppress a decrease of capacity. In addition, the type of battery is not limited thereto since

it is easy to attach the negative electrode tab to the metallic lithium plate.

The negative electrode for a lithium battery according to the present invention can be employed in any lithium battery. Particularly, it can be employed to a lithium-sulfur battery having a positive active material of a sulfuric material. The lithium-sulfur battery comprises: a negative electrode according to any one of the first to the fourth embodiments; a positive electrode comprising a positive active material selected from the group consisting of elemental sulfur,  $\text{Li}_2\text{S}_n$  ( $n \geq 1$ ),  $\text{Li}_2\text{S}_n$  ( $n \geq 1$ ) dissolved in a catholyte, an organosulfur compound, and a carbon-sulfur polymer  $((\text{C}_2\text{S}_x)_n$ ;  $x = 2.5$  to  $50$ ,  $n \geq 2$ ); and an electrolyte.

The electrolyte may be either a solid electrolyte or a liquid electrolyte.

The solid electrolyte can function as both a separator and a medium capable of transporting metal ions, and it can be composed of any ionic conductive material that is electrochemically stable. The ionic conductive material may include a glass electrolyte, a polymer electrolyte, or a ceramic electrolyte. The preferable solid electrolyte is made by adding an appropriate supporting electrolyte to a polymer electrolyte such as polyether, polyimine, polythioether, and so on. The solid electrolyte separator may comprise less than about 20% by weight of a non-aqueous organic solvent. In this case, it can further comprise a suitable gelling agent to reduce the fluidity of the organic solvent.

When the electrolyte is a liquid electrolyte, the lithium-sulfur battery should



further comprise a separator composed of porous glass, plastic, ceramic, or a polymer in order to physically separate the electrodes. The liquid electrode comprises a non-aqueous organic solvent and an electrolyte salt. The organic solvent may include a commonly used non-aqueous organic electrolyte such as ethylenecarbonate, propylenecarbonate, dioxolane, sulfolane, xylene, diglyme, tetrahydrofuran, tetraglyme, and so on.

The electrolyte salt may include a lithium cation-consisting lithium salt, an organic cation-consisting salt, or a mixture thereof.

The example of a lithium salt may include, but is not limited to,  $\text{LiPF}_6$ ,  $\text{LiBF}_4$ ,  $\text{LiSbF}_6$ ,  $\text{LiAsF}_6$ ,  $\text{LiClO}_4$ ,  $\text{LiCF}_3\text{SO}_3$ ,  $\text{Li}(\text{CF}_3\text{SO}_2)_2\text{N}$ ,  $\text{LiC}_4\text{F}_9\text{SO}_3$ ,  $\text{LiSbF}_6$ ,  $\text{LiAlO}_4$ ,  $\text{LiAlCl}_4$ ,  $\text{LiN}(\text{C}_x\text{F}_{2x+1}\text{SO}_2)(\text{C}_y\text{F}_{2y+1}\text{SO}_2)$  (wherein  $x$  and  $y$  are natural numbers),  $\text{LiCl}$ ,  $\text{LiI}$ , and so on.

The organic cation-consisting salt has a low vapor pressure, a very high flash point, and anti-combustibility, rendering the battery safe and anti-corrosive, and so that it is formed as a mechanically stable film. The preferable salt may include a large organic cation having a van der Waals volume of more than  $100^3$ . The greater the van der Waals volume of such a cation, the less the lattice energy, so the ion conductivity is reduced.

The organic cation-consisting salt can be present as a liquid phase in a wide range of temperatures. The organic cation-consisting salt is preferably

present as a liquid phase at a temperature of less than 100 °C, more preferably present as a liquid phase at a temperature of less than 50 °C, and most preferably present as a liquid phase at a temperature of less than 25°C. It is to be understood that it can be present as a liquid phase at a different range of temperatures depending on the applied method.

The organic cation is preferably any cation of a heterocyclic compound. The hetero atom of the heterocyclic compound may be selected from the group consisting of N, O, S, or a combination thereof. The heterocyclic composition may have one to four hetero atoms, and preferably one or two hetero atoms. The cation of the heterocyclic compound includes a cation of the compound selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, and triazolium, or a substitute thereof. It is preferable a cation of an imidazolium compound such as 1-ethyl-3-methylimidazolium (EMI), 1,2-dimethyl-3-propylimidazolium (DMPI), 1-butyl-3-methylimidazolium (BMI), and so on.

An anion to be bound with the cation may be any one among bis(perfluoroethylsulfonyl)imide ( $(\text{N}(\text{C}_2\text{F}_5\text{SO}_2)_2^-$ , Beti), bis(trifluoromethylsulfonyl)imide ( $(\text{N}(\text{CF}_3\text{SO}_2)_2^-$ , Im), tris(trifluoromethylsulfonylmethide ( $\text{C}(\text{CF}_3\text{SO}_2)_2^-$ , Me), trifluoromethane sulfonamide, trifluoromethane sulfonimide, trifluoromethyl sulfonimide, trifluoromethyl sulfonate,

$\text{AsF}_6^-$ ,  $\text{ClO}_4^-$ ,  $\text{PF}_6^-$ ,  $\text{BF}_4^-$ , and so on.

The preferable example of the organic cation-containing salt includes 1-ethyl-3-methylimidazolium bis(perfluoroethyl sulfonyl)imide (EMIBeti), 1,2-dimethyl-3-propylimidazolium bis(trifluoromethyl sulfonyl)imide (DMPIm), or 1-butyl-3-methylimidazolium hexafluorophosphate (BMIPF<sub>6</sub>).

Hereinafter, the present invention will be explained in detail with reference to examples. These examples, however, should not in any sense be interpreted as limiting the scope of the present invention.

#### Example 1

75 wt.% of sulfur powder as a positive active material, 12 wt.% of polyethylene oxide (PEO) as a binder, and 13 wt.% of ketjen black as a conductive material were added to and uniformly dispersed with acetonitrile to prepare a positive active material slurry. The uniformly dispersed slurry was coated on a carbon-coated Al foil using a doctor blade to provide a positive electrode. Then, the positive electrode was cut into a size of 22 cm<sup>2</sup> and Al tabs were welded thereto to prepare a positive electrode plate.

200 μm thick lithium metal foil was cut into a size of 3X3 cm<sup>2</sup> and a part of the foil was rubbed three times with a brush in order to impart a roughness to the surface thereof. The average surface roughness of the metallic lithium foil was 0.5 μm, determined using an optical 3D profiling system (Model No. NT2000, fabricated by WYKO). 10 μm thick copper foil was positioned on the surface of

the lithium metal foil treated to have the average surface roughness, and it was then pressed with a pressure of about 0.3 tonnes to obtain a negative electrode plate.

The obtained positive electrode plate, a vacuum-dried polyethylene separator, and the obtained negative electrode plate were subsequently stacked and inserted into a pouch. An electrolyte of 1M  $\text{LiN}(\text{CF}_3\text{SO}_2)$  in 1,3-dioxolane/dimethoxyethane/diglyme (2:4:4 volume ratio) was injected therein, and the pouch was sealed to complete a pouch-type test-cell.

#### Example 2

A test cell was fabricated by the same procedure as described in Example 1 except that the negative electrode plate was fabricated in a manner of cutting 200  $\mu\text{m}$  thick lithium metal foil to a size of 3X3  $\text{cm}^2$ , stacking a 100  $\mu\text{m}$  thick nickel foam having 85% porosity on the lithium foil, and pressing them at about 0.3 tonnes.

#### Example 3

A test cell was fabricated by the same procedure as described in Example 1 except that the negative electrode plate was fabricated in a manner of cutting 200  $\mu\text{m}$  thick lithium metal foil into a size of 3X3  $\text{cm}^2$ , stacking 10  $\mu\text{m}$  thick copper foils on both surfaces of the lithium foil, and welding them.

#### Comparative Example 1

A test cell was fabricated by the same procedure as described in Example 1 except that the negative electrode plate was fabricated in a manner of cutting 200  $\mu\text{m}$  thick lithium metal foil into a size of 3X3  $\text{cm}^2$ , stacking 100  $\mu\text{m}$  thick nickel tab on the lithium foil, and pressing them.

30 test cells for each of Examples 1 to 3 and Comparative Example 1 were fabricated and the internal resistance (IR) and open circuit voltage (OCV) were measured. The results are shown in Table 1. IR and OCV were determined using a model 3550 (fabricated by HIOKI E.E. Corporation).

【Table 1】

	Example 1		Example 2		Example 3		Comparative Example 1	
	IR( $\Omega$ )	OCV	IR( $\Omega$ )	OCV	IR( $\Omega$ )	OCV	IR( $\Omega$ )	OCV
1	5.4	3.20	9.3	3.23	8.3	3.22	off	3.20
2	5.8	3.21	8.9	3.19	9.0	3.22	24	3.24
3	5.5	3.20	8.5	3.21	9.7	3.20	26	3.20
4	5.3	3.20	8.8	3.22	16.0	3.22	27	3.20
5	4.8	3.20	10.9	3.04	7.8	3.22	15	3.22
6	5	3.20	10.0	3.17	11.4	3.20	off	3.13

7	5.5	3.22	9.2	3.12	16.0	3.21	25	3.23
8	5.5	3.20	11.7	3.18	15.0	3.23	23	3.22
9	5.3	3.20	6.8	3.22	10.0	3.26	27	3.17
10	6.2	3.20	10.5	3.21	9.6	3.25	off	3.18
11	4.3	3.20	12.5	3.17	9.2	3.16	23	3.19
12	4.0	3.20	10.3	3.21	9.5	3.16	22	3.19
13	3.9	3.20	15.0	3.22	11.6	3.18	17	3.19
14	4.5	3.20	11.1	3.20	10.7	3.14	25	3.21
15	2.8	3.20	7.7	3.22	9.9	3.12	off	3.17
16	4.6	3.20	7.0	3.22	12.4	3.18	off	3.17
17	4.7	3.20	11.3	3.20	7.5	3.22	off	3.21
18	4.2	3.20	8.5	3.21	11.2	3.21	29	3.20
19	4.0	3.21	9.2	3.23	13.2	3.17	24	3.24
20	4.2	3.20	15.0	3.26	11.0	3.21	22	3.20
21	5.2	3.20	7.3	3.25	9.6	3.19	26	3.20
22	4.5	3.20	10.9	3.16	12.5	3.23	24	3.22
23	4.3	3.20	15.5	3.23	11.3	3.19	28	3.18
24	4.2	3.20	14.5	3.26	12.0	3.21	29	3.15
25	4.8	3.20	9.5	3.25	8.6	3.22	off	3.22

26	4.5	3.20	9.1	3.16	7.9	3.04	25	3.17
27	4.8	3.20	8.7	3.16	12.2	3.17	26	3.18
28	4.7	3.20	9.0	3.18	9.4	3.17	23	3.18
29	5.2	3.20	11.1	3.18	9.5	3.16	off	3.21
30	4.8	3.20	10.2	3.18	9.0	3.12	off	3.20

Note: the indication “off” means the internal resistance is more than 30  $\Omega$ .

As shown in Table 1, the internal resistance of test cells of Examples 1 to 3 according to the present invention was significantly lower than those of Comparative Example 1. Since an increased internal resistance indicates an unstable contact between the tab and the electrode, it is advantageous to have the low internal resistance of the test cells of Examples 1 to 3 of the present invention, indicating a stable contact between the negative electrode and the negative electrode tab.

#### **【EFFECT OF THE INVENTION】**

The negative electrode for a lithium battery according to the present invention can reduce the internal resistance upon charge and discharge of the battery due to the firm attachment between the metallic lithium plate and the negative electrode tab. A decrease of capacity is thereby prevented by

decreasing the internal resistance so that it is possible to provide a high capacity battery. Further, it facilitates attaching the negative electrode tab to the metal lithium plate so that the type of the battery to be fabricated is not limited, and the occurrence of shorts is reduced.

While the present invention has been described in detail with reference to the preferred embodiments, those skilled in the art will appreciate that various modifications and substitutions can be made thereto without departing from the spirit and scope of the present invention as set forth in the appended claims.



## 【CLAIMS】

1. A negative electrode for a lithium battery, comprising:  
a metallic lithium plate; and  
a negative tab attached to the surface of the metallic lithium plate,  
wherein the average surface roughness of the metallic lithium plate on the  
area attached to the negative tab is 0.1 to 5  $\mu\text{m}$ .
2. The negative electrode for a lithium battery according to claim 1, wherein  
the average surface roughness is 0.3 to 0.6  $\mu\text{m}$ .
3. The negative electrode for a lithium battery according to claim 1, wherein  
the metallic lithium plate is a metallic lithium foil or metallic lithium coated on a  
conductive substrate.
4. The negative electrode for a lithium battery according to claim 3, wherein  
the conductive substrate is selected from the group consisting of a metal foil, a  
metal film, a conductive polymer film, and a polymer film deposited with a metal.
5. The negative electrode for a lithium battery according to claim 1, wherein  
the negative electrode tab is a 10 to 50  $\mu\text{m}$  thick metal plate or metal foam.
6. The negative electrode for a lithium battery according to claim 1, wherein  
the negative electrode tab is made of at least one metal selected from the group

consisting of nickel, copper, iron, and stainless steel.

7. A negative electrode for a lithium battery, comprising:  
a metallic lithium plate; and  
a negative electrode tab attached to the metallic lithium plate,  
wherein the negative electrode tab has a porosity of 50 to 100%.
8. The negative electrode for a lithium battery according to claim 7, wherein the metallic lithium plate is a metallic lithium foil or metallic lithium coated on a conductive substrate.
9. The negative electrode for a lithium battery according to claim 8, wherein the conductive substrate is selected from the group consisting of a metal foil, a metal film, a conductive polymer film, and a polymer film deposited with a metal.
10. The negative electrode for a lithium battery according to claim 7, wherein the negative electrode tab is composed of at least one metal selected from the group consisting of nickel, copper, iron, and stainless steel.
11. The negative electrode for a lithium battery according to claim 7, wherein the porosity of the negative electrode tab is 80 to 95%.
12. A negative electrode for a lithium battery, comprising:  
a metallic lithium plate; and  
a negative electrode tab attached to both the upper and lower end surfaces of the metallic lithium plate.

13. The negative electrode for a lithium battery according to claim 12, wherein the metallic lithium plate is a metallic lithium foil or metallic lithium coated on a conductive substrate.

14. The negative electrode for a lithium battery according to claim 13, wherein the conductive substrate is selected from the group consisting of a metal foil, a metal film, a conductive polymer film, and a polymer film deposited with a metal.

15. The negative electrode for a lithium battery according to claim 13, wherein the negative electrode tab is made of at least one metal selected from the group consisting of nickel, copper, iron, and stainless steel.

16. A method of fabricating a negative electrode for a lithium battery, comprising the steps of:

brushing the surface area of a metallic lithium plate to be attached to a negative electrode tab so that the average surface roughness of the surface area (Ra) is 0.1 to 5  $\mu\text{m}$ ; and

pressing the negative electrode tab onto the metallic lithium plate and attaching the negative electrode tab to the metallic lithium plate.

17. A lithium battery comprising a negative electrode according to any one of claims 1 to 15.

18. A lithium-sulfur battery comprising:

a negative electrode according to claims 1 to 15;

a positive electrode comprising a positive active material selected from the group consisting of elemental sulfur,  $\text{Li}_2\text{S}_n$  ( $n \geq 1$ ),  $\text{Li}_2\text{S}_n$  ( $n \geq 1$ ) dissolved in a catholyte, an organosulfur compound, and a carbon-sulfur polymer  $((\text{C}_2\text{S}_x)_n$ :  $x = 2.5$  to  $50$ ,  $n \geq 2$ ); and

an electrolyte.

【DRAWINGS】

【Fig 1】

